

Serial No. 09/960,449  
Filed September 21, 2001  
Amendment

Amendments to the Claims

Claims 1-4, 8-11, 13-17, 21-23, and 25 are pending in the application. Claims 1 and 14 have been amended to add the element that the pendant crosslinkable groups are acrylamide groups containing olefinically unsaturated groups. The amendment is supported in the specification on page 8, lines 22-24. Please amend claims 1 and 14 as follows:

1. (currently amended) A hydrogel wound dressing formed by spray delivery of a liquid composition to the wound, wherein the composition comprises water soluble PVA macromers having one or more pendant crosslinkable groups and the macromers crosslink to form a hydrogel *in situ* on the wound, wherein the pendant crosslinkable groups are acrylamide groups containing olefinically unsaturated groups.

14. (currently amended) A method of forming a hydrogel wound dressing, comprising the step of applying a composition to a wound via spray, wherein the composition comprises water soluble PVA macromers having one or more pendant crosslinkable groups and the macromers crosslink to form a hydrogel on the wound, wherein the pendant crosslinkable groups are acrylamide groups containing olefinically unsaturated groups.

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Listing of Claims

1. (currently amended) A hydrogel wound dressing formed by spray delivery of a liquid composition to the wound, wherein the composition comprises water soluble PVA macromers having one or more pendant crosslinkable groups and the macromers crosslink to form a hydrogel *in situ* on the wound, wherein the pendant crosslinkable groups are acrylamide groups containing olefinically unsaturated groups.
2. (original) The wound dressing of claim 1, wherein the hydrogel is degradable.
3. (original) The wound dressing of claim 1, wherein the composition is delivered via an aerosol delivery device.
4. (original) The wound dressing of claim 1, wherein the composition is delivered via a pump spray delivery device.
- 5-7. (cancelled)
8. (original) The wound dressing of claim 1, wherein the composition further contains one or more additives selected from the group consisting of preservatives, biologically active agents, defoamers, wetting agents, leveling agents, hydrating agents, thickeners, fillers, and absorbents.
9. (previously presented) The wound dressing of claim 8, wherein the active agent is selected from the group consisting of growth factors, nitric oxide, antibiotics, anti-inflammatories, analgesics, blood coagulants, and enzymes.
10. (original) The wound dressing of claim 8, wherein the active agent is one which delivers NO to the wound.
11. (original) The wound dressing of claim 1, wherein the dressing debrides the wound when it is removed.
12. (cancelled)
13. (previously presented) The wound dressing of claim 1, wherein the *in situ* crosslinking is in response to redox initiation.
14. (currently amended) A method of forming a hydrogel wound dressing, comprising the step of applying a composition to a wound via spray, wherein the composition comprises water soluble PVA macromers having one or more pendant crosslinkable groups and

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the macromers crosslink to form a hydrogel on the wound, wherein the pendant crosslinkable groups are acrylamide groups containing olefinically unsaturated groups.

15. (original) The method of claim 14, wherein the hydrogel is degradable.
16. (original) The method of claim 14, wherein the composition is delivered via an aerosol delivery device.
17. (original) The method of claim 14, wherein the composition is delivered via a pump spray delivery device.
- 18- 20. (cancelled)
21. (original) The method of claim 14, wherein the composition further contains one or more additives selected from the group consisting of preservatives, biologically active agents, defoamers, wetting agents, leveling agents, hydrating agents, thickeners, fillers, and absorbents.
22. (previously presented) The method of claim 21, wherein the active agent is selected from the group consisting of growth factors, nitric oxide, antibiotics, anti-inflammatories, analgesics, blood coagulants, and enzymes.
23. (original) The method of claim 21, wherein the active agent is one which delivers NO.
24. (cancelled)
25. (previously presented) The method of claim 14, wherein the *in situ* crosslinking is in response to redox initiation.
26. (cancelled)

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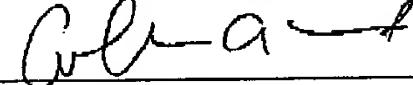
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Conclusion

Reconsideration of the claims as amended is respectfully requested.

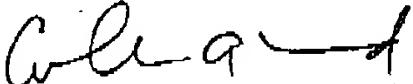
Respectfully submitted,

  
Collen A. Beard  
Registration No. 38,824

Law Office of Collen A. Beard, LLC  
P.O. Box 1064  
Decatur, Georgia 30031-1064  
(404)373.5065

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